Protective humoral and CD4⁺ T cellular immune responses of *Staphylococcus aureus* vaccine MntC in a murine peritonitis model

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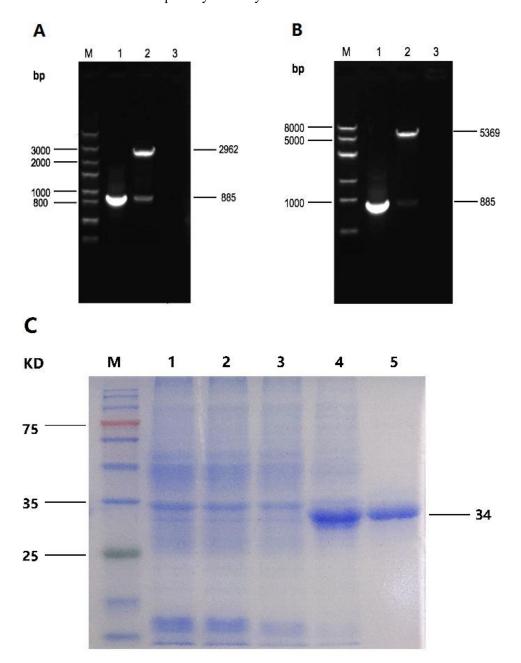
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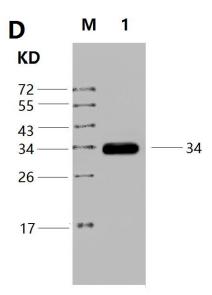
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Supplementary Information

This file contains 2 supplementary figures.

Supplementary Figure S1. Cloning, expression and purification of recombinant MntC. (A) Identification of MntC-pMD18T by PCR and double enzyme digestion. Lane M: DNA Marker DL5000; Lane 1: PCR product; Lane 2: pMD18T-MntC; Lane 3: Negative control; (B) Identification of MntC-pET28a by PCR and double enzyme digestion. Lane M: DNA Marker DL8000; Lane 1: PCR product; Lane2: pET28a-MntC; Lane3: Negative control; (C) SDS-PAGE and western-blotting analysis of expressed and purified recombinant protein MntC. Lane M: PAGE Ruler Prestained Protein Ladder; Lane 1: E. coli BL21(DE3) with pET-28a before induction; Lane 2: E. coli BL21 (DE3) with pET-28a induced by IPTG; Lane 3: E. coli BL21 (DE3) with pET-28a/MntC before induction; Lane 4: E. coli BL21 (DE3) with pET-28a/MntC induced by IPTG; Lane 5: Purified MntC protein. (D) Western blotting of purified MntC, anti-His antibodies were tested for primary antibody.





Supplementary Figure S2. Purity of isolated CD4⁺ **T cells.** Isolated CD4⁺ **T cells treated with** anti-CD4-PE demonstrated by flow cytometry analyses, the purity was greater than 95%.

